

Day : Friday
Date: 1/14/2005
Time: 16:03:41

PALM INTRANET

Inventor Name Search Result

Your Search was:

Last Name = GORGOJO LOBATO

First Name = JOSE

Application#	Patent#	Status	Date Filed	Title	Inventor Name 1
10810128	Not Issued	030	03/26/2004	PROCESS FOR OBTAINING 17BETA-(SUBSTITUTED)-3-OXO-DELTA 1,2-4-AZASTEROIDS AND INTERMEDIATES	GORGOJO LOBATO, JOSE MARIA

Inventor Search Completed: No Records to Display.

Search Another:

Inventor

Last Name

Gorgojo Lobato

First Name

Jose

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STN- Structure Search
1.14.05

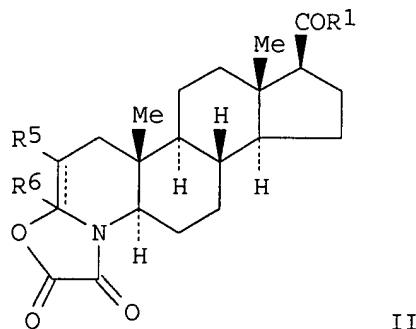
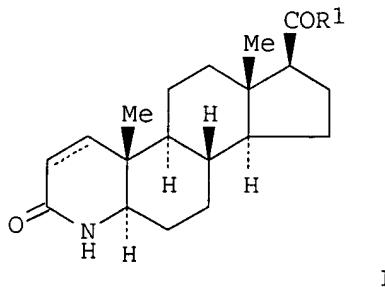
10/810,128

=> d ibib abs hitstr 1-2

inventor

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN
 ACCESSION NUMBER: 2003:282583 CAPLUS
 DOCUMENT NUMBER: 138:287866
 TITLE: Process for the preparation of 17 β -substituted-3-oxo- Δ 1,2-4-azasteroids and intermediates thereof
 INVENTOR(S): Gorgojo Lobato, Jose Maria; Lorente Bonde-Larsen, Antonio; Martin Juarez, Jorge
 PATENT ASSIGNEE(S): Ragactives, S.L., Spain
 SOURCE: PCT Int. Appl., 27 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Spanish
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003029267	A2	20030410	WO 2002-ES453	20020926
WO 2003029267	A3	20030619		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
ES 2185503	A1	20030416	ES 2001-2190	20010929
ES 2185503	B1	20040801		
EP 1437361	A2	20040714	EP 2002-779579	20020926
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
US 2004254209	A1	20041216	US 2004-810128	20040326
PRIORITY APPLN. INFO.:			ES 2001-2190	A 20010929
			WO 2002-ES453	W 20020926
OTHER SOURCE(S): GI		CASREACT 138:287866; MARPAT 138:287866		



AB The present invention discloses a process for preparing 17 β -substituted-3-oxo- Δ 1,2-4-azasteroids, such as I [R1 = alkyl, OR2; R2 = alkyl, NR3R4; R3, R4 = H, alkyl; dashed line = double bond], from 17 β -substituted-3-oxo-4-azasteroids I [dashed line = single bond].

Thus, I [R1 = NHBu-t; dashed line = single bond] was reacted with oxanyl chloride to provide oxazolidinedione derivative II [R1 = NHBu-t; R5,R6 = H; dashed line = double bond], which upon reaction with 1,3-dibromo-5,5-dimethyl-hydantoin in presence of perchloric acid afford 2-bromo-3-hydroxyoxazolididione derivative III [R1 = NHBu-t; R5 = Br, R6 = OH; dashed line = single bond (III)]. III was reacted with potassium tert-butoxide in presence of anhydrous DMF to afford I [R1 = NHBu-t; dashed line = double bond]. Some prepared compds. are inhibitors of testosterone-5 α -reductase and can be used in the treatment of hyperandrogenic alterations.

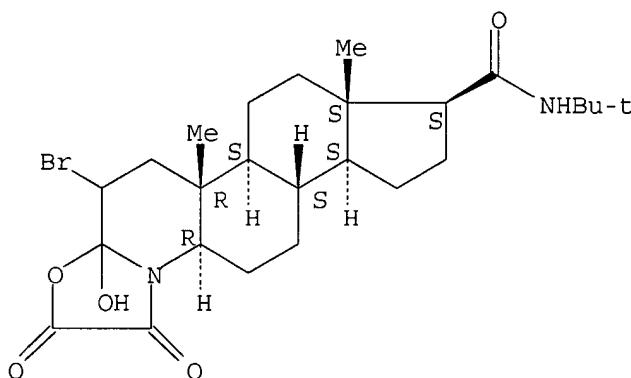
IT 135252-06-1P 507221-52-5P 507221-53-6P

RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of 17 β -substituted-3-oxo- Δ 1,2-4-azasteroids and intermediates thereof)

RN 135252-06-1 CAPPLUS

CN Indeno[5,4-f]oxazolo[3,2-a]quinoline-8-carboxamide, 4-bromo-N-(1,1-dimethylethyl)octadecahydro-3a-hydroxy-5a,7a-dimethyl-1,2-dioxo-, (5aR,5bS,7aS,8S,10aS,10bS,12aR)- (9CI) (CA INDEX NAME)

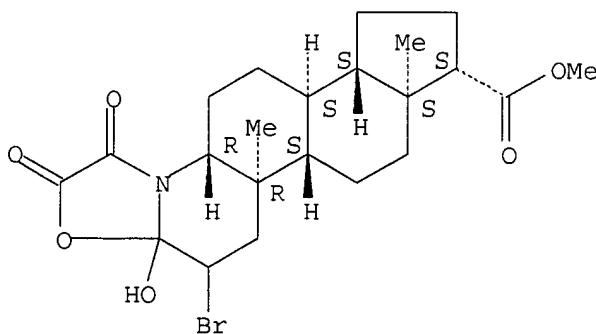
Absolute stereochemistry.



RN 507221-52-5 CAPPLUS

CN Indeno[5,4-f]oxazolo[3,2-a]quinoline-8-carboxylic acid, 4-bromo-octadecahydro-3a-hydroxy-5a,7a-dimethyl-1,2-dioxo-, methyl ester, (5aR,5bS,7aS,8S,10aS,10bS,12aR)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

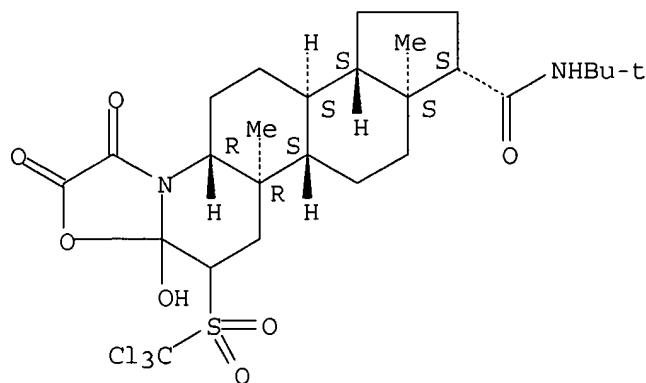


RN 507221-53-6 CAPPLUS

CN Indeno[5,4-f]oxazolo[3,2-a]quinoline-8-carboxamide, N-(1,1-dimethylethyl)octadecahydro-3a-hydroxy-5a,7a-dimethyl-1,2-dioxo-4-[(trichloromethyl)sulfonyl]-, (5aR,5bS,7aS,8S,10aS,10bS,12aR)- (9CI) (CA INDEX NAME)

INDEX NAME)

Absolute stereochemistry.



L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1991:472017 CAPLUS

DOCUMENT NUMBER: 115:72017

TITLE: Method for introducing a 1,2 double bond into azasteroids

INVENTOR(S) : King, Anthony O.; Weinstock, Leonard M.; Anderson, Kevin R.; Shuman, Richard F.

PATENT ASSIGNEE(S): Merck and Co., Inc., USA

SOURCE: Eur. Pat. Appl., 9 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

DOCUMENT TYPE: Recent
LANGUAGE: English

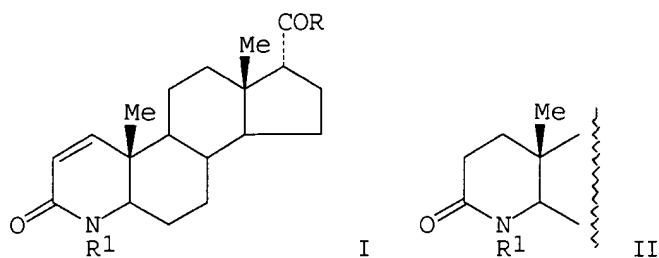
LANGUAGE.
FAMILY ACT.

FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT INFORMATION:

RECEIVED NO

EP 428366	A2	19910522	EP 1990-312341	19901113	
EP 428366	A3	19920729			
EP 428366	B1	19950920			
R: CH, DE, FR, GB, IT, LI, NL					
US 5021575	A	19910604	US 1989-434663	19891113	
CA 2029859	AA	19910514	CA 1990-2029859	19901113	
CA 2029859	C	20020514			
JP 03206096	A2	19910909	JP 1990-304208	19901113	
JP 06051718	B4	19940706			
EP 655459	A2	19950531	EP 1995-200326	19901113	
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EP 655459	B1	20000503			
R: CH, DE, FR, GB, IT, LI, NL					
LV 12572	B	20010420	LV 2000-117	20000907	
RITY APPLN. INFO.:				US 1989-434663	A 19891113
				EP 1990-312341	A3 19901113
R SOURCE(S):		CASREACT 115:72017; MARPAT 115:72017			



AB 1,2-Unsatd. azasteroids I [R = H, (un)substituted C1-12 alkyl, cycloalkyl, Ph, OH, alkoxy, OCH₂Ph, amino; R1 = H, Me, Et] were prepared from saturated derivs. II in a 3-step 1-pot reaction. Thus, II (R = CMe₃, R1 = H) was converted to oxazolidinedione derivs. with oxalyl chloride, brominated with Br, treated with MeNHCH₂CH₂OH to hydrolyze the oxazolidinedione, and dehydrobrominated with Me₃COK. The overall yield of I (R = CMe₃, R1 = H) was 60.2%.

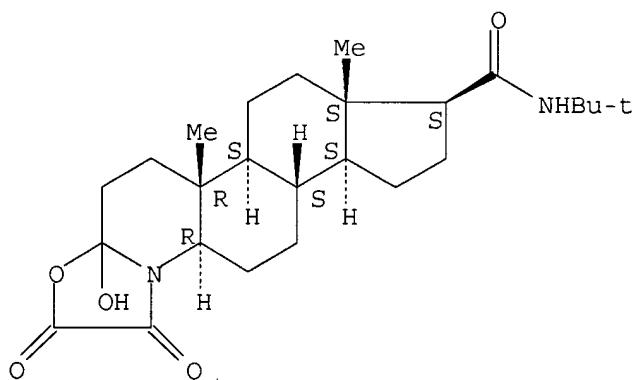
IT 135252-05-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and bromination of)

BN 135252-05-0 CAPIUS

RN 135232-03-0 C1425
CN 4-Azaandrostane-4-acetic acid, 17-[[$(1,1\text{-dimethylethyl})$ amino]carbonyl]-3,3-dihydroxy- α -oxo-, γ -lactone, ($5\alpha,17\beta$)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



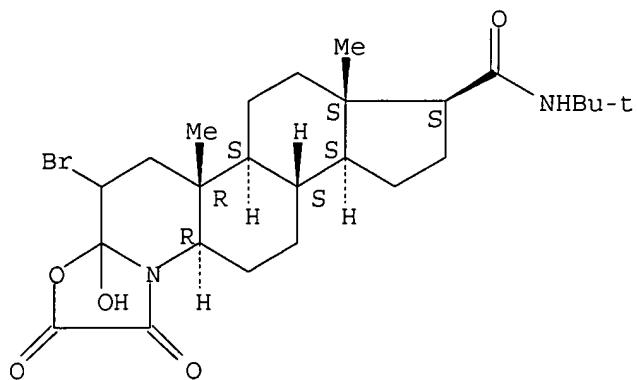
IT 135252-06-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and oxazolidinedione hydrolysis of)

RN 135252-06-1 CAPLUS

CN Indeno[5,4-f]oxazolo[3,2-a]quinoline-8-carboxamide, 4-bromo-N-(1,1-dimethylethyl)octadecahydro-3a-hydroxy-5a,7a-dimethyl-1,2-dioxo-, (5aR,5bS,7aS,8S,10aS,10bS,12aR)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



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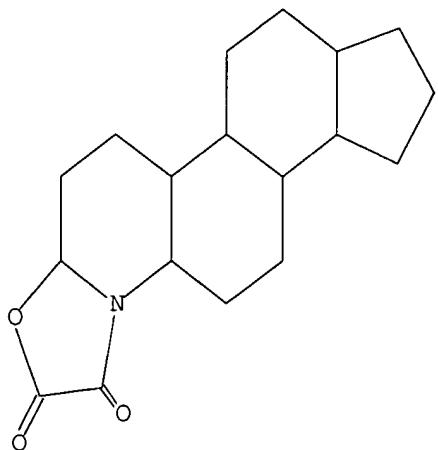
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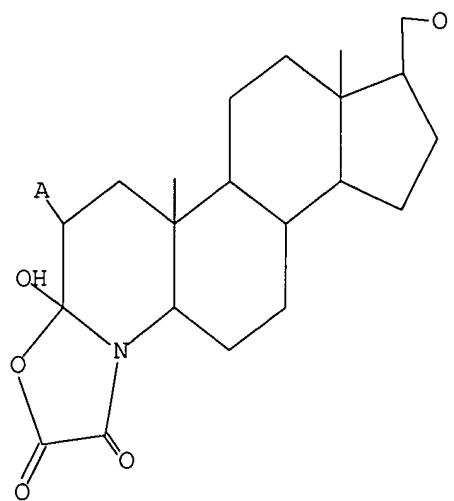
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L3 HAS NO ANSWERS
L3 STR



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=> d 11
L1 HAS NO ANSWERS
L1 STR

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Structure attributes must be viewed using STN Express query preparation.

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